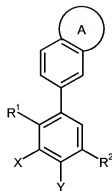


Amendments to the claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

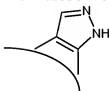
1. (Currently amended) A compound of formula (I):



(I)

wherein

A is a fused 5-membered heteroaryl ring selected from



or



, which ring is

substituted by $-(CH_2)_m$ heterocyclyl wherein the heterocyclyl is a 5- or 6-membered heterocyclic ring ~~containing one or two heteroatoms independently selected from tetrahydrofuranyl, tetrahydropyranlyl or morpholinyl-oxygen, sulfur, and nitrogen~~ optionally substituted by up to two substituents independently selected from oxo, C_{1-6} alkyl, $-(CH_2)_n$ phenyl, $-OR^3$, $-(CH_2)_nCO_2R^3$, $-NR^3R^4$ and $-CONR^3R^4$, and

A is optionally further substituted by one substituent selected from $-OR^3$, halogen, trifluoromethyl, $-CN$, $-CO_2R^3$ and C_{1-6} alkyl optionally substituted by hydroxy;

R^1 is selected from methyl and chloro;

R^2 is selected from $-NH-CO-R^5$ and $-CO-NH-(CH_2)_q-R^6$;

R^3 and R^4 are each independently selected from hydrogen and C_{1-6} alkyl;

R^5 is selected from hydrogen, C_{1-6} alkyl, $-(CH_2)_q-C_{3-7}$ cycloalkyl, trifluoromethyl, $-(CH_2)_r$ heteroaryl optionally substituted by R^7 and/or R^8 , and $-(CH_2)_r$ phenyl optionally substituted by R^7 and/or R^8 ;

R⁶ is selected from hydrogen, C₁₋₆alkyl, C₃₋₇cycloalkyl, -CONHR⁹, phenyl optionally substituted by R⁷ and/or R⁸, and heteroaryl optionally substituted by R⁷ and/or R⁸;

R⁷ is selected from C₁₋₆alkyl, C₁₋₆alkoxy, -(CH₂)_q-C₃₋₇cycloalkyl, -CONR⁹R¹⁰, -NHCOR¹⁰, halogen, -CN, -(CH₂)_sNR¹¹R¹², trifluoromethyl, phenyl optionally substituted by one or more R⁸ groups, and heteroaryl optionally substituted by one or more R⁸ groups;

R⁸ is selected from C₁₋₆alkyl, C₁₋₆alkoxy, halogen, trifluoromethyl, and -(CH₂)_sNR¹¹R¹²;

R⁹ and R¹⁰ are each independently selected from hydrogen and C₁₋₆alkyl, or

R⁹ and R¹⁰, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹³, wherein the ring may be substituted by up to two C₁₋₆alkyl groups;

R¹¹ is selected from hydrogen, C₁₋₆alkyl and -(CH₂)_q-C₃₋₇cycloalkyl optionally substituted by C₁₋₆alkyl,

R¹² is selected from hydrogen and C₁₋₆alkyl, or

R¹¹ and R¹², together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹³;

R¹³ is selected from hydrogen and methyl;

X and Y are each independently selected from hydrogen, methyl and halogen;

m and q are each independently selected from 0, 1 and 2;

n and r are each independently selected from 0 and 1; and

s is selected from 0, 1, 2 and 3;

with the proviso that:

A is not substituted by -(CH₂)_mNR¹⁴R¹⁵ wherein R¹⁴ and R¹⁵, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulphur and NR¹⁶ wherein R¹⁶ is hydrogen or methyl,

when m is 0, the -(CH₂)_mheterocyclyl group is not a 5- or 6-membered heterocyclyl ring containing nitrogen optionally substituted by C₁₋₂alkyl or -(CH₂)_nCO₂R³, and

the compound of formula (I) is not 1,1-dimethylethyl 4-(6-{5-[(cyclopropylamino)carbonyl]-2-methylphenyl}-1,2-benzisoxazol-3-yl)-1-piperazinecarboxylate;

or a pharmaceutically acceptable [[derivative]] salt thereof.

2. (Currently amended) A compound according to claim 1 wherein A is a ~~fused-5-membered heteroaryl ring containing two heteroatoms independently selected from oxygen and nitrogen~~ substituted by the $-(CH_2)_m$ heterocyclyl moiety on the ring nitrogen of the A ring.

3. (Currently amended) A compound according to claim 1 wherein A is substituted by $-(CH_2)_m$ heterocyclyl and [[wherein]] the heterocyclyl is an optionally substituted tetrahydropyranyl a 5- or 6-membered ring containing one or two heteroatoms independently selected from oxygen and nitrogen optionally substituted by up to two substituents independently selected from oxo, C_{1-6} alkyl, $-(CH_2)_n$ phenyl, $-OR^3$, $-(CH_2)_nCO_2R^3$, $-NR^3R^4$, and $-CONR^3R^4$.

4. (Previously Presented) A compound according to claim 1 wherein R^1 is methyl.

5. (Previously Presented) A compound according to claim 1 wherein R^2 is $-CO-NH-(CH_2)_q-R^6$.

6. (Previously Presented) A compound according to claim 1 wherein X is fluorine.

7. (Currently amended) A compound according to claim 1 ~~substantially as hereinbefore defined with reference to any one of Examples 1 to 9, which is~~
N-Cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2H-pyran-2-ylmethyl)-1H-indazol-5-yl]benzamide ;;

N-Cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2-furanylmethyl)-1H-indazol-5-yl]benzamide ;

3-{1-[(4-Benzylmorpholin-2-yl)methyl]-1H-indazol-5-yl}-*N*-cyclopropyl-5-fluoro-4-methylbenzamide ;

N-Ethyl-4-methyl-3-[3-(tetrahydro-3-furanyl)-1H-indazol-6-yl]benzamide ;

N-Ethyl-3-fluoro-4-methyl-5-[3-(tetrahydro-3-furanyl)-1H-indazol-6-yl]benzamide ;

or a pharmaceutically acceptable [[derivative]] salt thereof.

8. (Currently amended) A compound selected from:

N-cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2H-pyran-2-ylmethyl)-1H-indazol-5-yl]benzamide;

N-cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2-furanylmethyl)-1H-indazol-5-yl]benzamide; and

3-{1-[(4-benzylmorpholin-2-yl)methyl]-1H-indazol-5-yl}-*N*-cyclopropyl-5-fluoro-4-methylbenzamide,

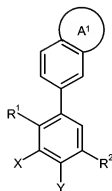
or a pharmaceutically acceptable [[derivative]] salt thereof.

9. (Previously Presented) A pharmaceutical composition comprising at least one compound as claimed in claim 1, or a pharmaceutically acceptable derivative thereof, in association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.

10 -13 (Cancelled)

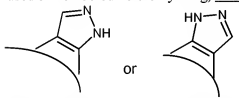
14. (Currently amended/Withdrawn) A process for preparing a compound of formula (I) as ~~claimed in~~ according to claim 1, or a pharmaceutically acceptable ~~[[derivative]]~~ salt thereof, which comprises:

(a) reacting a compound of formula (II)



(II)

in which R¹, R², X and Y are as defined in claim 1 and A¹ is ~~an unsubstituted fused 5-membered heteroaryl ring~~. A is a fused 5-membered heteroaryl ring selected from



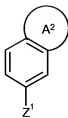
with a halide derivative of formula (III)



(III)

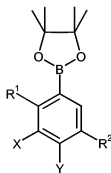
in which $-(CH_2)_m \text{heterocyclyl}$ is as defined in claim 1 and Z is halogen, in the presence of a base;

(b) reacting a compound of formula (IV)

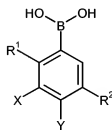


(IV)

in which A^2 is A as defined in claim 1 or a protected form of A or A^1 , and Z^1 is halogen, with a compound of formula (VA) or (VB)



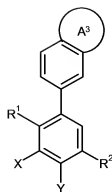
(VA)



(VB)

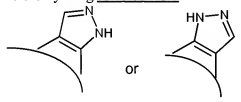
in which R^1 , R^2 , X and Y are as defined in claim 1, in the presence of a catalyst;

(c) reacting a compound of formula (XI)



(XI)

in which R^1 , R^2 , X and Y are as defined in claim 1 and A^3 is a fused 5-membered heteroaryl ring selected from



substituted by $-(CH_2)_m$ heterocyclyl wherein the heterocyclyl is unsubstituted, with a suitable reagent; or

(d) final stage modification of one compound of formula (I) as defined in claim 1 to give another compound of formula (I) as defined in claim 1.

15. (Currently amended) A compound according to claim [[2]] 1 wherein A is substituted by $-(CH_2)_m$ heterocyclyl and [[wherein]] the heterocyclyl is an optionally substituted tetrahydrofuranyl ~~a 5- or 6-membered ring containing one or two heteroatoms independently selected from oxygen and nitrogen optionally substituted by up to two substituents independently selected from oxo, C₁₋₆alkyl, $-(CH_2)_n$ phenyl, $-OR^3$, $-(CH_2)_nCO_2R^3$, $-NR^3R^4$, and $-CONR^3R^4$.~~

16. (Previously Presented) A compound according to claim 15 wherein R^1 is methyl.

17. (Previously Presented) A compound according to claim 15 wherein R^2 is $-CO-NH-(CH_2)_qR^6$.

18. (Previously Presented) A compound according to claim 15 wherein X is fluorine.

19. (new) A compound according to claim 1 wherein A is substituted by $-(CH_2)_m$ heterocyclyl and the heterocyclyl is an optionally substituted morpholinyl.
20. (new) A compound according to claim 3 wherein m is 1.
21. (new) A compound according to claim 15 wherein m is 0.
22. (new) A compound according to claim 15 wherein m is 1.
23. (new) A compound according to claim 19 wherein m is 1.
24. (new) A compound according to claim 1 wherein A is substituted by the $-(CH_2)_m$ heterocyclyl moiety on the 3-position of the A ring.
25. (new) A pharmaceutical composition comprising a compound according to claim 8, or a pharmaceutically acceptable salt thereof, in association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.